


NORTHROP EXHIBIT P

Cut
Remb

M. All. / 

47

Notes (Signal back of several (not all)
results photos with this pen (other (on front)
was ~~was~~ permanent ink)


- PCR (HIV - MSP) worked well in
integrated-heater device, gel electrophoresis
verified product. Some, but minimal Primase
(esp. due to known fact that device
react in mixture cycled 1-2 times,
then at R.T. for $\frac{1}{2}$ hr & prior to
20 cycles due to need to re-solder
connections - new rxn mixture (30 μ l)
was added)

- was able to extract $\sim 100\%$ of aqueous
phase with 200 μ l (set at 30 μ l)
pipette & load 5-6 wells of
electrophoresis channel
- (*) \rightarrow calculate power consumed in today's
experiment compare to batteries

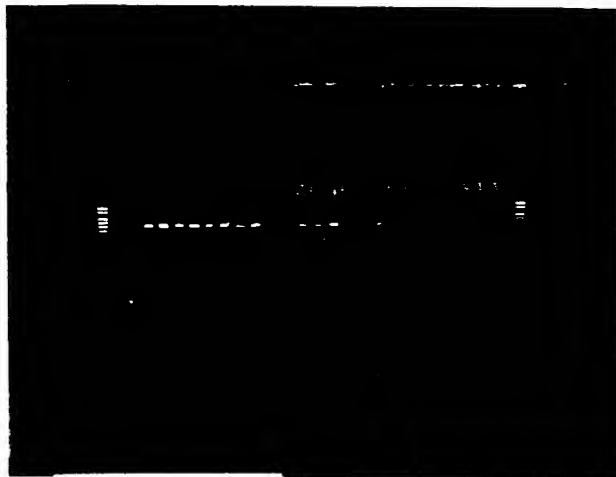
Other Discussion

- Last Tues w/ Ray Manilla
here (Cetus) along w/
Ross Higuchi, Bob Watson, Russ's
technician, myself we tried
homogeneous detection w/ video
CCD over 460 thermal cycles
- pulsed He-Ne laser (ILEE laser
company, Switz) was tried
 - \rightarrow see LLNH Book (notebook)
for details

Cont
Results (photos)
Devia PCB results positive

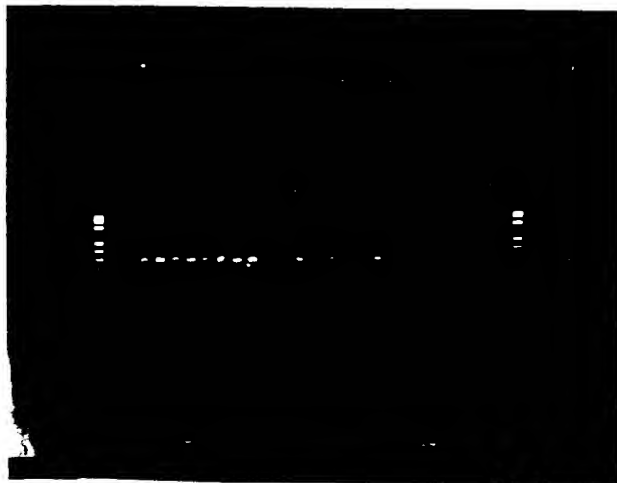
M. All ¹⁵ 

electr. T = 15 min




M. All  T = 24°C
4.6 3200

elect = 40 min



M. All  T = 24°C
4.6 3200

taped recipe from
Watson
K

Ren	50 pl 10x RM	2
pa		
B. Watson	50 pl 1mm d. 17p	3
	50 pl M13	4
	10 pl 10x10 = 100 pencils 10x10	5
M. Allen / 	10 pl photo	6
	2 1/2 pl 10x1.750/rev 12.5	
	12.5/50 pl = 2.5	
	→ 2.5	
	327.5 420	9
	<hr/>	
	500	

Cellus M. All ~~for~~

Try new PCR system
(more Temp forgiving)

142 bp product target as SS M13 from
gag-region of HIV

- 1) Starting target = 10^8 copies in 5 μ l
 $T = 96-55$ \downarrow 16-18 cycles
 (works at 88+) is plenty
- 2) primers

old names:		new names	
SK145	=	ph07	10 μ g/ μ l
SK431	=	ph08	

Reaction mixture: (500 μ l)

50 μ l 10x Buffer w/ MgCl
 " 1 mM dNTPs
 " M13 w/ gag region of HIV
 10 μ l = $10 \times 10 = 100$ pmols

10 μ l (same for) $\frac{\text{ph07}}{\text{ph08}}$?

2 1/2 μ l = $10 \times 1.25 \mu$ /pml 12.5
 Tag

327.5 μ l H₂O

500 μ l total rxn volume

500.0
 -172 1/2

 327.5

M. All ~~page~~

(cont)

- 1) re-use voltage (same device) as
on mail 30 (ie 3.17 V at 98°C)
at 0.2A

Do only 20 cycles

A) Standards

10, 10, 20, 20, 30, 30, 40, 40

-150 μ l oil (1-8)B) Device 30 μ l w ~ 90 μ l oil

1-minute cycles at 3.17V
20-1 minute cycles (A-E) 0.2A

Electrophoresis

well-problem

10 (10)

P Std 6 1 2 3 4 5 6 7 8 9 10 A B C D D E 1 2 3 4

- 1c) Had to re-solder device ^{wire connectors} after 2-cycles
fix time \approx 1/2 hour rxn was
at Room temp

Results - ① failed product in both
stds and in wells
② wells (and 1) std had
less bright primer - dimers
③ dev. needed ~6-5 μ l gel

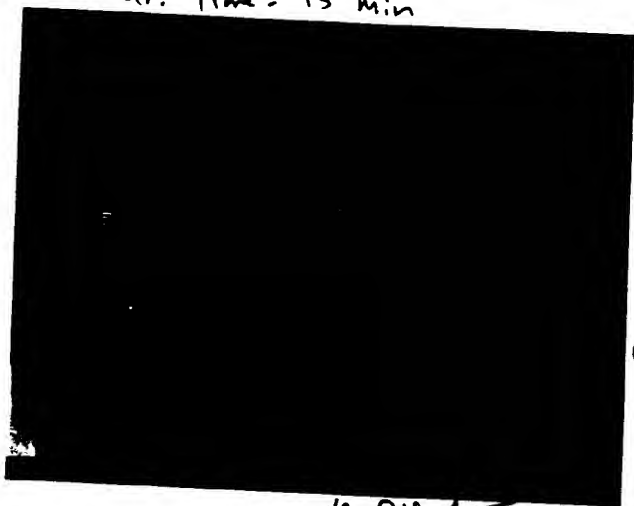
See next
2 pages:

44

Cont results (photos)

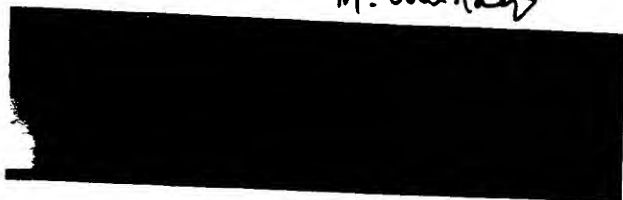
M. Allen

electr. Time = 15 min



T = 1 sec 5.6 3200

M. Allen



M

T = 1 sec 5.6 3200

*1 Loaned to Mila Ching